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FILE 'CAPLUS' ENTERED AT 12:58:32 ON 03 JUL 2003
L1 31 S NANOSOL
L2 27 S NANO SOL
L3 57 S L1 OR L2
L4 89738 S DRUG DELIVERY
L5 14539 S CHITOSAN
L6 16869 S AMPHIPHIL?
L7 140522 S COLLOID?
L8 256685 S L4 OR L5 OR L6 OR L7
L9 10 S L3 AND L8
L10 47 S L3 NOT L9
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ANSWER 1 OF 10 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2002:815104 CAPLUS

DOCUMENT NUMBER: 138:325012

TITLE: Deposition of Titania Nanoparticles on Spherical

Silica AUTHOR (S):

Ryu, Dong Hwan; Kim, Seong Chul; Koo, Sang Man; Kim,

Dong Pyo

CORPORATE SOURCE: College of Engineering, Ceramic Processing Research Center, Department of Chemical Engineering, Hanyang

University, Seoul, 133-791, S. Korea SOURCE:

Journal of Sol-Gel Science and Technology (2003),

26(1/2/3), 489-493

CODEN: JSGTEC; ISSN: 0928-0707 Kluwer Academic Publishers

DOCUMENT TYPE:

PUBLISHER:

Journal

LANGUAGE: English

Titania-coated silica nanoparticles were prepd. through a sol-gel process using peptized TiO2 nano-sols. The TiO2 sols were obtained by peptization, the process of redispersing a coagulated colloid, and were coated on SiO2 particles by the control of the wt. ratio of TiO2/SiO2 and the pH of the mixt. in aq. soln. At pH 4.5 the difference of zeta-potential between SiO2 and TiO2 maximized and then the TiO2-coated SiO2 particles with highest TiO2 contents (.apprx.20%) were obtained without the self-aggregation of TiO2 sols. The morphologies of particles were characterized with field emission SEM (FE-SEM) and TEM and the isoelec. points (IEP) of particles were measured by zeta potential. The nature of exposed titanium ions on the surface of titania-coated silica nanoparticles was elucidated by XPS.

REFERENCE COUNT: THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS 6 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 2 OF 10 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER:

DOCUMENT NUMBER:

2002:388038 CAPLUS 137:116835

TITLE:

Exfoliation and restacking route to anatase-layered titanate nanohybrid with enhanced photocatalytic

activity

AUTHOR (S):

Choy, Jin-Ho; Lee, Hyun-Cheol; Jung, Hyun; Kim,

Hasuck; Boo, Hankil

CORPORATE SOURCE:

National Nanohybrid Materials Laboratory and School of Chemistry and Molecular Engineering, Seoul National

University, Seoul, 151-747, S. Korea

SOURCE:

Chemistry of Materials (2002), 14(6), 2486-2491

CODEN: CMATEX; ISSN: 0897-4756

PUBLISHER: DOCUMENT TYPE: American Chemical Society

LANGUAGE:

Journal

English A new microporous TiO2-pillared layered titanate has been prepd. by

hybridizing the exfoliated titanate with the anatase TiO2 nanosol The stable colloidal nano-sheet was obtained by intercalating tetrabutylamine into the layered protonic titanate, HxTi2-x/4.box.x/404 . times. H2O (x = 0.67), with a lepidocrocite-like structure. The colloidal suspension of exfoliated titanate sheets was mixed with the monodispersed anatase TiO2 nanosol soln. prepd. by the hydrolysis of titanium isopropoxide with acetylacetone. The obtained nanohybrid was heated at 300 .degree.C for 2 h in order to complete the grafting reaction of intercalated anatase TiO2 nanosol on the interlayer surface of layered titanate. According to the x-ray diffraction anal. and N2 adsorption-desorption isotherms, it was found that the TiO2-pillared layered titanate showed a pillar height of .apprx.2 nm, a high surface area of .apprx.460 m2/g, and a pore size of .apprx.0.95 nm, indicating the formation of a microporous pillar structure. Its photocatalytic activity was evaluated by measuring the total vol. of H2 gas evolved during the irradn. of the catalyst suspensions in water. The H2 gas evolution was found to increase from the layered titanate (cesium and protonic form) to the unsupported TiO2 (acac-TiO2) and the TiO2-pillared layered titanate, because the electron and hole recombination in the pillared system is thought to be effectively suppressed because of electron transfer between guest and host. A marked enhancement in the activity by ca. 40 times was obtained for TiO2-pillared layered titanate compared to pristine compds. such as layered titanate and anatase TiO2 nanosol when Pt (0.3 wt %) was doped on the surface of the sample.

REFERENCE COUNT:

THERE ARE 39 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 3 OF 10 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2001:673142 CAPLUS DOCUMENT NUMBER: 135:294428

TITLE: Relaxation dynamics of processes in colloidal zirconia nanosols. Dependence on excitation energy and temperature AUTHOR (S): Emeline, A. V.; Serpone, N. CORPORATE SOURCE: Department of Chemistry and Biochemistry, Concordia University, Montreal, QC, H3G 1M8, Can. Chemical Physics Letters (2001), 345(1,2), 105-110 SOURCE: CODEN: CHPLBC; ISSN: 0009-2614 PUBLISHER: Elsevier Science B.V. DOCUMENT TYPE: Journal LANGUAGE: English Process relaxation dynamics occurring in colloidal ZrO2 nanosols under steady-state and pulsed laser excitation are studied; the temp. dependence of the photoluminescence is also reported. The laser-induced emission emanating from these nanosols is thermally quenched with activation energy Etq = 0.048 .+-. 0.008 eV, identical to the activation energy (Etq = 0.050 .+-. 0.005 eV) of thermal quenching of the photoluminescence under steady-state excitation. It is also identical to the energy (Eads = 0.046 .+-. 0.005 eV) of thermal quenching of the photostimulated adsorption of O on ZrO2 micron-size particles. A pathway to summarize the events is proposed. THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 18 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 4 OF 10 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2001:614038 CAPLUS DOCUMENT NUMBER: 135:182238 TITLE: Composite of partially crystalline cellulose and modified silica gel as a carrier INVENTOR (S): Suess, Wolfgang; Boettcher, Horst; Kallies, Karl-Heinz PATENT ASSIGNEE(S): Feinchemie G.m.b.H. Sebnitz, Germany SOURCE: Ger. Offen., 5 pp. CODEN: GWXXBX DOCUMENT TYPE: Patent LANGUAGE: German FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. DATE -----A1 20010823 DE 10006125 DE 2000-10006125 20000211 PRIORITY APPLN. INFO.: DE 2000-10006125 20000211 A modified microcryst. cellulose (MC) comprising a composite of partially cryst. cellulose (a cellulose hydrolyzate with d.p. 30-400) and modified silica gel is suitable for use as a carrier for immobilization of colorants, indicators, pharmaceutical agents or biol. active substances in the pharmaceutical, cosmetics and food industries, biotechnol. and genetic engineering, as an auxiliary agent for pharmaceutical and cosmetic formulations, as a stabilizer of suspensions and thermally stable oil-in-water emulsions, as well as for use as a chromatog. support. preferably in column chromatog., a filtering material and an adsorbent. The SiO2 is modified with RSiOn, R2SiOn, A12O3, ZrO2 or TiO2, in which R =H, alkyl, aryl, epoxyalkyl or aminoalkyl and n = 1.5 or 1. The process for manufg. the modified MC comprises (1) impregnation of the suspended MC with the corresponding modified SiO2 nanosol, which is obtained by alk. or acid hydrolysis of tetraalkoxysilanes, optionally contg. substituted alkoxysilanes or metal alcoholates, (2) optionally addn. of solid or dissolved colorants, indicators, pharmaceutical agents or biol. active substances, (3) gelation by neutralization, addn. of fluoride, warming and evapn. of the solvent, and (4) drying, milling and sieving. Thus, to 9.46 g Al2(OH)5Cl.2.5H2O, dissolved in 86 mL H2O and 344 mL EtOH, 80.41 g Si(OEt) 4 was added under stirring within 10 min and stirred at room temp. for 24 h. Then, to 192.3 g of the aluminosilicate (.apprx.5.2% SiO2-Al2O3 in 85% EtOH) 6 mL 0.1% NH4OH soln. was added and 490 g microcryst. cellulose was impregnated, which was dried 30 min at 60.degree.. The composite obtained had a higher breaking strength, compressibility, and flowability than com. modified microcryst. cellulose.

THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 5 OF 10 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2001:470578 CAPLUS

DOCUMENT NUMBER: 135:197530

REFERENCE COUNT:

TITLE: Hydroxyapatite nano sol prepared via a mechanochemical route AUTHOR (S):

3

Nakamura, S.; Isobe, T.; Senna, M. CORPORATE SOURCE: Faculty of Science and Technology, Keio University,

SOURCE: Journal of Nanoparticle Research (2001), 3(1), 57-61

CODEN: JNARFA; ISSN: 1388-0764

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PUBLISHER:
                            Kluwer Academic Publishers
  DOCUMENT TYPE:
                            Journal
  LANGUAGE:
                            English
       Well-dispersed sol with cryst. hydroxyapatite was obtained directly by
       milling a mixt. comprising Ca(OH)2, an ag. soln. of H3PO4 and a
       dispersant, an ammonium salt of polyacrylic acid. The av. crystallite
       size of hydroxyapatite was below 20 nm. Ca/P molar ratio of the product
       was 1.51 .+-. 0.04, i.e., Ca deficient from stoichiometric hydroxyapatite.
       Min. apparent viscosity was attained at a dispersant concn. 0.92 wt.% of
       sol. An as-milled sol was dild. by a factor 2.61-10 solid wt.% to give a Newtonian fluid of 2 mPa.cntdot.s. From the dild. sol, we obtained a few
       .mu.m thick dense film of hydroxyapatite by dip coating on the slide glass
       precoated by chitosan.
  REFERENCE COUNT:
                                  THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS
                                  RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
      ANSWER 6 OF 10 CAPLUS COPYRIGHT 2003 ACS
  ACCESSION NUMBER:
                           2001:150590 CAPLUS
  DOCUMENT NUMBER:
                           134:183520
  TITLE:
                           Pharmaceuticals containing chitosan
                           derivative
  INVENTOR (S):
                           Hoffmann, Hans-Rainer; Asmussen, Bodo
  PATENT ASSIGNEE(S):
                           Lts Lohmann Therapie-Systeme A.-G., Germany
 SOURCE:
                           Ger. Offen., 4 pp.
                           CODEN: GWXXBX
 DOCUMENT TYPE:
                           Patent
 LANGUAGE:
                           German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:
      PATENT NO.
                       KIND DATE
                                              APPLICATION NO. DATE
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      DE 19940794
                      A1 20010301
                                             DE 1999-19940794 19990827
      WO 2001015669
                        A1 20010308
                                             WO 2000-EP7904 20000814
          W: AU, BR, CA, CN, CZ, HU, IL, IN, JP, KR, MX, NZ, PL, RU, TR, US, ZA RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
              PT, SE
      EP 1206253
                        A1 20020522
                                              EP 2000-964016 20000814
          R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY
      BR 2000013563 A
                              20020709
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                                                               20000814
      NZ 517541
                        Α
                              20020828
                                             NZ 2000-517541
                                                               20000814
                           20030304
      JP 2003508423
                        T2
                                             JP 2001-519883
                                                               20000814
                                          DE 1999-19940794 A 19990827
 PRIORITY APPLN. INFO.:
                                          WO 2000-EP7904 W 20000814
      Solid pharmaceuticals contain a partly loaded drug in nanosol
      form, in which the drug is bound to a charged chitosan deriv.
      Procedures for their prodn. and their use in the manuf. of pharmaceuticals
      are described.
     ANSWER 7 OF 10 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER:
                          1999:329503 CAPLUS
DOCUMENT NUMBER:
                          131:50050
TITLE:
                          Entrapment of nanostructured palladium clusters in
                          hydrophobic sol-gel materials
AUTHOR (S):
                          Reetz, Manfred T.; Dugal, Markus
CORPORATE SOURCE:
                          Max-Planck-Institut fur Kohlenforschung, Mulheim an
                          der Ruhr, D-45470, Germany
SOURCE:
                          Catalysis Letters (1999), 58(4), 207-212
                          CODEN: CALEER; ISSN: 1011-372X
PUBLISHER:
                          Baltzer Science Publishers
DOCUMENT TYPE:
                          Journal
LANGUAGE:
                          English
     The fluoride-catalyzed hydrolysis of mixts. of MeSi(OMe)3 and Mg(OEt)2 in
     the presence of preformed nanosized R4N+Br--stabilized Pd colloids
     gave micro/mesoporous hydrophobic sol-gel materials in which the Pd
     clusters are entrapped individually in the solid matrix. The materials
     are active catalysts in the hydrogenation of 1,5-cyclooctadiene.
REFERENCE COUNT:
                                THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS
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                                RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 8 OF 10 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER:
                         1998:335033 CAPLUS
DOCUMENT NUMBER:
                         129:28901
TITLE:
                         Polysiloxane-mineral fiber composites
INVENTOR (S):
                         Jonschker, Gerhard; Mennig, Martin; Schmidt, Helmut;
                         Angenendt, Rainer
PATENT ASSIGNEE(S):
                         Institut fuer neue Materialien gemeinnuetzige G.m.b.H.
                         Universitaet des Saar, Germany; Pfleiderer
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Daemmstofftechnik International G.m.b.H. und Co.

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SOURCE:

Ger. Offen., 6 pp. CODEN: GWXXBX

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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PATENT NO.
                        KIND DATE
                                                APPLICATION NO. DATE
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      DE 19647369
                               19980520
                                                DE 1996-19647369 19961115
      EP 842967
                         A2 19980520
                                                EP 1997-120034 19971114
                         A3
      EP 842967
                               19980617
                        B1
      EP 842967
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              IE, SI, LT, LV, FI, RO
      WO 9821266
                        A1 19980522
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                                                                  19971114
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              PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG,
         US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA,
              GN, ML, MR, NE, SN, TD, TG
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                        A1 19980603
B2 20010802
                                                AU 1998-56533
                                                                  19971114
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                         A 19991026
A 19991201
     BR 9712766
                                                BR 1997-12766
                                                                  19971114
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     ES 2150731
                         T3 20001201
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                                               JP 1998-522192
                                                                  19971114
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                         C2 20020910
                                               RU 1999-112565
                                                                  19971114
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                         Α
                              20000825
                                               KR 1999-703533
                                                                  19990422
     MX 9904099
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                              20001130
                                               MX 1999-4099
                                                                  19990430
     US 6187426
                         B1 20010213
                                               US 1999-297574
                                                                  19990503
PRIORITY APPLN. INFO.:
                                            DE 1996-19647369 A 19961115
                                            WO 1997-EP6371 W 19971114
```

The title composites comprise glass, mineral, or woody fibers in functional contact with nanocomposites prepd. by modifying the surface of colloidal, inorg. particles with silanes of specified structure under the conditions of the sol-gel process to form nanosols. Stirring a 65:15:20 MeSi(OEt)3-PhSi(OEt)3-Si(OEt)4 mixt. with a SiO2 sol and HCl formed a 14% polysiloxane nanosol which, after .apprx.12 h, was dild. to 0.5 mol H2O/alkoxy group, sprayed on moistened glass wool, and cured at 200 degree. for 5-10 min to give an elastic composite with better fire resistance than a phenolic resin-glass composite.

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ANSWER 9 OF 10 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER:
                       1998:335032 CAPLUS
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1

DOCUMENT NUMBER:

129:28900

TITLE:

Polysiloxane nanocomposites

INVENTOR(S): PATENT ASSIGNEE(S): Jonschker, Gerhard; Mennig, Martin; Schmidt, Helmut Institut fuer neue Materialien gemeinnuetzige G.m.b.H.

Universitaet des Saar, Germany

SOURCE:

Ger. Offen., 7 pp. CODEN: GWXXBX

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

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PATENT NO.
                         KIND DATE
                                                        APPLICATION NO. DATE
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DE 19647368
                         A1
                                 19980520
                                                        DE 1996-19647368 19961115
WO 9822648
                         A2 19980528
                                                        WO 1997-EP6370 19971114
WO 9822648
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      W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ,
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           PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US,
     PT, KU, KU, SD, SE, SG, SI, SA, SB, IU, IE, IK, II, OA, OS, OS, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG
WO 9822536
                         A2 19980528
A3 19980730
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WO 9822536
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               GN, ML, MR, NE, SN, TD, TG
      AU 9854825
                        A1 19980610
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      EP 946313
                        A2 19991006
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      EP 950039
                        A2 19991020
                                             EP 1997-951901
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          R: AT, BE, CH, DE, ES, FR, GB, IT, LI, NL, SE, PT, IE, FI
      CN 1236339
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PRIORITY APPLN. INFO.:
                                          DE 1996-19647368 A 19961115
                                          WO 1997-EP6370 W 19971114
                                          WO 1997-EP6372
                                                         W 19971114
                                          WO 1997-EP6373
                                                          W 19971114
     Composites are prepd. by hydrolyzing silanes of specified structure in the
     presence of colloidal inorg. particles under the conditions of
     the sol-gel process to form nanosols which, after further
     hydrolysis-condensation, are cured in contact with substrates. Stirring a
     60:20:20 MeSi(OEt)3-Si(OEt)4-PhSi(OEt)3 mixt. strongly with a SiO2 sol and
     HCl gave a sol which, after 12 h at 60.degree., was dild. with H20 to 0.5
     mol H20/alkoxy group and mixed with sand (particle size .apprx.1 mm) to
     give a sand content of 84% and cured at 100.degree. for 20 min to give a
     mech. stable composite which did not lose its shape after 1 h at
     500.degree..
    ANSWER 10 OF 10 CAPLUS COPYRIGHT 2003 ACS
L9
ACCESSION NUMBER:
                         1993:456207 CAPLUS
DOCUMENT NUMBER:
                          119:56207
TITLE:
                         Nanosol drug formulation
INVENTOR (S):
                         Wunderlich, Jens Christian; Schick, Ursula; Werry,
                         Juergen; Freidenreich, Juergen
PATENT ASSIGNEE(S):
                         Alfatec-Pharma GmbH, Germany
SOURCE:
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PCT Int. Appl., 72 pp.
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        German
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
    PATENT NO.
                     KIND DATE
                                          APPLICATION NO. DATE
    WO 9310768
                     A1
                          19930610
                                          WO 1992-DE1010 19921204
        W: AU, CA, JP, US
        RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
    DE 4140177
                     A1 19930609
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DE 1991-4140178 19911205

DE 1991-4140195 19911205

19921204

AU 1992-30802

DE 4140177

DE 4140178

DE 4140178

DE 4140195

DE 4140195

AU 9230802

C2

A1

C2

A1

C2

19951221

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19941027

19930628

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AU 671965
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            EP 615445
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US 5932245
PRIORITY APPLN. INFO.:
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                                                                 19990803
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DE 1991-4140178 A 19911205
                                                                                             DE 1991-4140195 A 19911205
WO 1992-DE1010 A 19921204
          Colloidally-dispersed solns. of drugs with low soly. in water (
          Colloidally-dispersed solns. of drugs with low soly. In water (nanosols) are prepd. by isoionic stabilization of the drug particles with gelatin, gelatin derivs. or collagen hydrolyzates of opposite charge. The nanosols show little Ostwald maturation and provide rapid drug absorption and high bioavailability. An ibuprofen nanosol was prepd. using a gelatin B (isoelec. point 4.9) which has a pos. net charge at pH 3. A 3% aq. gelatin B soln. (500 g) was adjusted to pH 3 and treated with 250 mL EtOH and 10 g ibuprofen. EtOH evaps. led to a nanosol, which was spray-dried.
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evapn. led to a nanosol, which was spray-dried.

L10 ANSWER 1 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2003:472121 CAPLUS TITLE: Nonlinear optical absorption in silver nanosol AUTHOR (S): Unnikrishnan, K. P.; Nampoori, V. P. N.; Ramakrishnan, V.; Umadevi, M.; Vallabhan, C. P. G. CORPORATE SOURCE: International School of Photonics, Cochin University of Science and Technology, Cochin, 682022, India SOURCE: Journal of Physics D: Applied Physics (2003), 36(11), 1242-1245 CODEN: JPAPBE; ISSN: 0022-3727 PUBLISHER: Institute of Physics Publishing DOCUMENT TYPE: Journal LANGUAGE: English Nonlinear optical absorption in silver nanosol was investigated at selected wavelengths (456 nm, 477 nm and 532 nm) using open aperture Z-scan technique. It was obsd. that nature of nonlinear absorption is sensitively dependent on input fluence as well as on excitation wavelength. Besides, the present sample was found to exhibit reverse saturable absorption (RSA) and saturable absorption (SA) at these wavelengths depending on excitation fluence. RSA is attributed to enhanced absorption resulting from photochem. changes. SA obsd. for fluence values lower and higher than those corresponding to RSA are, resp., attributed to plasmon bleach and satn. of RSA. L10 ANSWER 2 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2003:427272 CAPLUS TITLE: Efficient transdermal penetration and improved stability of L-ascorbic acid encapsulated in an inorganic nanocapsule AUTHOR (S): Yang, Jae-Hun; Lee, Sun-Young; Han, Yang-Su; Park, Kyoung-Chan; Choy, Jin-Ho CORPORATE SOURCE: National Nanohybrid Materials Laboratory, School of Chemistry & Molecular Engineering, Seoul National University, Seoul, 151-747, S. Korea Bulletin of the Korean Chemical Society (2003), 24(4), SOURCE: 499-503 CODEN: BKCSDE; ISSN: 0253-2964 PUBLISHER: Korean Chemical Society DOCUMENT TYPE: Journal LANGUAGE: English Encapsulation of L-ascorbic acid (vitamin C) within a bio-compatible layered inorg. material was achieved by copptn. reaction, in which the layered inorg. lattice and its intercalate of vitamin C are simultaneously formed. The nano-meter sized powders of vitamin C intercalate thus prepd. was again encapsulated with silica nano-sol to form a nanoporous shell structure. This ternary nanohybrid of vitamin C-layered inorg. core-SiO2 shell exhibited an enhanced storage stability and a sustained releasing of vitamin C. Furthermore, the nano-encapsulation of vitamin C with inorg. mineral was very helpful in delivering vitamin C mols. into skin through stratum corneum, facilitating transdermal penetration of vitamin C in topical application. REFERENCE COUNT: THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS 26 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L10 ANSWER 3 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2003:355803 CAPLUS DOCUMENT NUMBER: 138:342352 TITLE: Sol-gel preparation of low phonon energy silicate glass-ceramic gain medium with SnO2 nanoclusters for lasers and optical amplifiers INVENTOR(S): Taylor, Elizabeth; Brambilla, Gilberto; Chiodini, Norberto; Paleari, Alberto; Spinolo, Giorgio; Morazzoni, Franca; Scotti, Roberto PATENT ASSIGNEE(S): UK SOURCE: U.S. Pat. Appl. Publ., 13 pp. CODEN: USXXCO DOCUMENT TYPE: Patent LANGUAGE: English FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. DATE ------US 2003087742 A1 20030508 US 2002-260414 20021001 PRIORITY APPLN. INFO.: EP 2001-308392 A 20011002 US 2001-327768P P 20011010 Optical gain medium are fabricated by sol-gel process and include a glass ceramic host material contg. clusters of cryst. oxide material (such as SnO2) and doped with active ions (such as rare earth, erbium or thulium)

concd. at the clusters. The active ions are preferentially located at the nanoclusters so that they experience the relatively low phonon energy of the oxide and are insensitive to the phonon energy of the host. A host with a high phonon energy, such as silica, can therefore be used without the usual drawback of reduced carrier lifetimes through enhanced nonradiative decay rates.

L10 ANSWER 4 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2003:317353 CAPLUS

DOCUMENT NUMBER: 138:325943

TITLE: Bio-composite for the removal of heavy metals from

aqueous solutions

INVENTOR (S): Kallies, Karl-Heinz; Selenska-Pobell, Sonja; Raff,

Johannes; Soltmann, Ulrich; Boettcher, Horst; Quast,

PATENT ASSIGNEE(S): Kallies Feinchemie AG, Germany; Forschungszentrum

Rossendorf E.V. SOURCE: Ger. Offen., 6 pp. CODEN: GWXXBX

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO KIND DATE APPLICATION NO. DATE DE 10146375 A1 20030424 DE 2001-10146375 20010920 PRIORITY APPLN. INFO.:

DE 2001-10146375 20010920 A bio-composite material for the biosorption of heavy metals from aq. solns. consists of drying-stable cell products which are homogeneously dispersed in an inorg. gel. The inorg. gel can contain oxides of group IIA-VA and IIB-VB elements. The inorg. gel is a SiO2 nanosol prepd. by hydrolysis of tetraalkoxysilanes and can be modified by Co-hydrolysis with metal alcoholates or halides, tri- or dialkoxysilanes. The cell product, such as spores, cell wall proteins or killed cells, are mixed with the aq. nanosol. A gel is formed by layer formation, neutralization, heating and/or fluoride addn. The gel is freeze-dried or air-dried.

L10 ANSWER 5 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2003:208013 CAPLUS

TITLE . Studies on nano-particle sols of hydroxyapatite and

titanium dioxide for haemo-compatibility AUTHOR (S): Chen, Xiao; Feng, Lingyun; Peng, Renxiu; Cao,

Xianying; Li, Shipu Department of Pharmacology, Medical School of Wuhan CORPORATE SOURCE:

University, Wuhan, 430071, Peop. Rep. China

Weisheng Yanjiu (2002), 31(3), 197-199 CODEN: WEYAEM; ISSN: 1000-8020

PUBLISHER: Weisheng Yanjiu Bianjibu

DOCUMENT TYPE: Journal

LANGUAGE · Chinese

SOURCE:

A biol. evaluation was conducted for two types of nano- particle sols, hydroxyapatite(HAP) and titanium dioxide(TiO2). The results showed that HAP sol significantly prolonged the bleeding time and coagulation time of mice as well as the prothrombin time(PT) and partial thromboplastin time(PTT) of rats while TiO2 sol exhibited no such effects. Neither HAP sol nor TiO2 sol instigated in vitro hemolysis of rabbit erythrocyte. However, both of the materials caused in vitro aggregation of rabbit erythrocytes . The reason underlying the different results as to the two types of material was their specific stablizer, heparin for HAP sol and PVC for TiO2 sol. A biol. inert stablizer has no less significance than the nano-particle's very own nature in a name material's application prospect.

L10 ANSWER 6 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2003:205181 CAPLUS

TITLE: Al2O3-Y-TZP/Al2O3 functionally graded composites of

tubular shape from nano-sols using double-step electrophoretic deposition

AUTHOR (S): Kaya, Cenqiz

CORPORATE SOURCE: Interdisciplinary Research Centre (IRC) in Materials

Processing and School of Metallurgy and Materials, The University of Birmingham, Edgbaston, Birmingham, B15

2TT

SOURCE: Journal of the European Ceramic Society (2003),

23(10), 1655-1660 CODEN: JECSER; ISSN: 0955-2219

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English

Al203-Y-TZP/Al203 functionally graded composites of tubular shape incorporating a very tough central layer with graded compn. (Al2O3-Y-TZP) and a hard outer surface layer of pure alumina were produced from nano-size sols using electrophoretic deposition (EPD) in an attempt to generate a continuously inhomogeneous property variation across the final component and to control the microstructure at a nanometer scale. It is shown that hardness, fracture toughness and alumina grain size within the graded layer are controlled by the vol. fraction of TZP grains and the highest vol. fraction (71%) of TZP phase provides a fracture toughness value of 7.1 MPa m1/2 and Vicker's hardness of 10.4 GPa while the lowest vol. fraction (13%) results in obtaining a fracture toughness value of 3.8 MPa m1/2 and hardness value of 15.7 GPa. The pure alumina surface layer (100 .mu.m in thickness) with a high hardness value of 19.4 GPa is considered to be beneficial for tribol. applications where high wear resistance is required.

L10 ANSWER 7 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2003:132147 CAPLUS TITLE:

Structural evolution of SiO2-ZrO2 nano-

sol intercalated clays upon pillaring reaction AUTHOR (S): Choy, Jin-Ho; Yoon, Joo-Byoung; Jung, Hyun; Park,

Joo-Hyoung

CORPORATE SOURCE: National Nanohybrid Materials Laboratory, School of

Chemistry & Molecular Engineering, Seoul National

University, Seoul, 151-747, S. Korea SOURCE:

Journal of Materials Chemistry (2003), 13(3), 557-562

CODEN: JMACEP; ISSN: 0959-9428 Royal Society of Chemistry

DOCUMENT TYPE: Journal

PUBLISHER:

LANGUAGE: English

A two-dimensional layered nanohybrid with a high sp. surface area has been prepd. by ion exchange reactions between the sodium ions in montmorillonite and the pos. charged Zr-coated SiO2 sol particles. The basal spacing increases from 12.5 .ANG. to 26 .ANG. upon intercalation due to the insertion of a SiO2-ZrO2 nano-sol into the interlayer space of montmorillonite. Upon calcining at 300 .degree.C, it transforms into a porous nanohybrid with a basal spacing of 22.6 .ANG. The N2 adsorption-desorption isotherms were characterized as being of type IV according to the BDDT classification, indicating the existence of a large no. of micro- and mesopores. From its hysteresis curves, one can classify the nanohybrid as being of the H3 type with slit-shaped pores by the IUPAC classification. The estd. BET sp. surface area and av. micropore size are about 358 m2 g-1 and 12 .ANG., resp., with the latter value is similar to the gallery height of the sample indicating that the SiO2-ZrO2 sol particles are intercalated to form a monolayer. The local structural evolution of the Zr species in SiO2-ZrO2 sol particle has been investigated systematically by X-ray absorption spectroscopy at the Zr K-edge with respect to the calcination temp. The EXAFS spectroscopic results confirm that the surface of the nano-sized SiO2 sol particles is coated with the Zr species.

REFERENCE COUNT: 28 THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 8 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2002:922173 CAPLUS

DOCUMENT NUMBER: 138:126339 TITLE:

Biosorption of uranium and copper by biocers AUTHOR (S):

Raff, J.; Soltmann, U.; Matys, S.; Selenska-Pobell,

S.; Boettcher, H.; Pompe, W.

CORPORATE SOURCE: Institute of Radiochemistry, Research Center

Rossendorf, Dresden, D-01314, Germany

Chemistry of Materials (2003), 15(1), 240-244 SOURCE:

CODEN: CMATEX; ISSN: 0897-4756

PUBLISHER: American Chemical Society DOCUMENT TYPE:

Journal LANGUAGE: English

Biol. ceramic composites (biocers) made according to an aq. sol-gel protocol were used as selective metal binding filters. The biol. component of the biocers, Bacillus sphaericus JG-A12, was isolated from a uranium mining waste pile. Vegetative cells and spores of this strain are known to selectively bind U, Cu, Al, Cd, and Pb in large amts. Sol-gel ceramics were prepd. by dispersing vegetative cells, spores, and stabilized surface-layer proteins (S-layer) in aq. silica nanosols gelling, and drying. The biosorption of uranium and copper by the three kinds of biocers and by their single components was investigated with regard to dependence on time, concn., and prepn. conditions. Biocers with cells possess the highest binding capacity as compared to matrixes with spores and an S-layer. Freeze-drying of prepd. biocers or adding water-sol. compds. such as sorbitol lead to higher porosity and faster

metal binding. Uranium was bound mainly to the biol. component but also to the SiO2 network. Copper was only bound by the cells, spores, or S-layer. Bound uranium and copper were completely removed by washing with aq. citric acid.

REFERENCE COUNT:

PUBLISHER:

THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 9 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2002:815241 CAPLUS

DOCUMENT NUMBER: 138:357724

TITLE: Biosorption of heavy metals by sol-gel immobilized Bacillus sphaericus cells, spores and S-layers Soltmann, U.; Raff, J.; Selenska-Pobell, S.; Matys, AUTHOR (S):

S.; Pompe, W.; Boettcher, H.

CORPORATE SOURCE: Forschungszentrum Rossendorf, Institut fuer

Radiochemie, Dresden, D-01314, Germany

SOURCE: Journal of Sol-Gel Science and Technology (2003),

26(1/2/3), 1209-1212

CODEN: JSGTEC; ISSN: 0928-0707 Kluwer Academic Publishers

DOCUMENT TYPE: Journal

LANGUAGE: English

Different types of biocers were prepd. by dispersing vegetative cells, spores and surface layer proteins (S-layers) of Bacillus sphaericus JG-A12 in aq. silica nanosols, gelling or depositing on glass, and drying. The enzymic activity of embedded B. sphaericus cells depends noticeably on the water content of the biocer. The cells are destroyed by drying and shrinkage of the silica network, whereas embedded spores retain their ability for germination. The biosorption of uranium and copper on these biocomposites was investigated. Biocers with cells possess the highest metal binding capacity compared to matrixes with spores or S-layers. An addnl. increment of the metal binding capacity is achieved by using penetration reagents like sorbitol. For renewed use biosorbed uranium and copper can be completely removed from the biocers by using aq. citric acid. The use of spores as biocomponent offers possibilities for the prepn. of storage-stable bioactive biocers.

REFERENCE COUNT: 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 10 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2002:675475 CAPLUS

DOCUMENT NUMBER: 137:359955

TITLE: Sol-Gel Derived Gold Nanoclusters in Silica Glass

Possessing Large Optical Nonlinearities

AUTHOR (S): Selvan, S. Tamil; Hayakawa, Tomokatsu; Nogami,

Masayuki; Kobayashi, Yoshio; Liz-Marzan, Luis M.;

Hamanaka, Yasushi; Nakamura, Arao

Department of Materials Science Engineering, Nagoya CORPORATE SOURCE:

Institute of Technology, Gokiso-cho, Showa-ku, Nagoya,

466-855, Japan

SOURCE: Journal of Physical Chemistry B (2002), 106(39).

10157-10162

CODEN: JPCBFK; ISSN: 1520-6106

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

Sol-gel derived Au/SiO2 glasses, starting with Au sols prepd. by 2 different methodologies, 1 without the aid of any external stabilizing agent and the other with SiO2-coated Au, are described. The optical absorption spectra show the typical surface plasmon resonance for Au at .apprx.520-530 nm. TEM reveals the existence of spherical Au particles in the SiO2 matrix. The mean diams. of Au nanoclusters in gels and glasses vary from 10 to 20 nm and from 27 to 38 nm, resp., supported by x-ray diffraction data. The 3rd-order optical nonlinearities |.chi.(3)| detd. by a degenerate 4-wave mixing (DFWM) method exhibit a higher value of 2.2 times. 10-9 esu for a glass with 0.5% Au heated at 600.degree. indicating that a larger size induces an enhancement effect. This method also offers a greater .chi.(3)/.alpha. (.alpha.: absorption coeff.) value of 1 .times. 10-11 esu cm, which is comparable to those values obtained by a sputtering method. Thus, these novel Au-SiO2 nanocomposites fabricated by facile sol-gel methods render them potential candidates for photonic applications.

REFERENCE COUNT: THERE ARE 54 CITED REFERENCES AVAILABLE FOR THIS 54 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 11 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2002:465198 CAPLUS

DOCUMENT NUMBER: 137:385924

TITLE: Refining of textiles by nanosol coating Mahltig, B.; Bottcher, H.

AUTHOR (S):

CORPORATE SOURCE: Gesellschaft zur Forderung von Medizin-, Bio- und Umwelttechnologien e.V., GMBU, Dresden, Germany Melliand Textilberichte (2002), 83(4), E50-E51, SOURCE . 251-253 CODEN: MTIRDL; ISSN: 0341-0781 PUBLISHER: Melliand Textilberichte DOCUMENT TYPE: Journal LANGUAGE: English The sol-gel technique is a new versatile tool to create transparent and well adhesive metal or silicon oxide films on various textiles. To synthesize such coatings on textiles, silica contg. nanosol suspensions were used to impregnate the textile materials. After impregnation a drying process converts the applied nanosol into a thin and transparent gel film. The chem. and phys. modification of such coatings allows a wide range in the variation of surface properties of textiles. REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L10 ANSWER 12 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2002:366930 CAPLUS DOCUMENT NUMBER: 136:352322 TITLE: Biocomposite material containing inorganic gel and live cells INVENTOR (S): Boettcher, D. Horst; Sawusch, Stefan; Kallies, Karl-Heinz; Pompe, Wolfgang PATENT ASSIGNEE(S): Feinchemie Gmbh Sebnitz, Germany SOURCE: Ger. Offen., 8 pp. CODEN: GWXXBX DOCUMENT TYPE: Patent LANGUAGE: German FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. DATE DE 10054119 A1 20020516 DE 2000-10054119 20001031 PRIORITY APPLN. INFO.: DE 2000-10054119 20001031 The invention concerns a biocomposite material and its prepn. from an inorg. gel and live cells, e.g. microorganisms; the biocomposite materials are used as biocatalysts. Culture medium, activators, e.g. org. acids can be included. Thus an aq. silica nanosol was prepd. from a tetrahydroxysilane ethanolic soln. with 0.01 M hydrochloric acid, neutralized with ammonia and dryed. Saccharomyces cerevisiae was dispersed in 1% Tween 80 - contg. water and the nanosol gel was added. Series compns. were prepd. with activator combinations, e.g. org. acid and EDTA, org. ammonium salt and tetra-Bu ammonium chloride, water-sol. org. compd. and sorbit. The fermentative properties of the biocatalyst products were tested with glucose; the activator-contg. products showed increased activity. REFERENCE COUNT: THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L10 ANSWER 13 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2002:298333 CAPLUS DOCUMENT NUMBER: 137:85821 TITLE: A new nanohybrid photocatalyst between anatase (TiO2) and layered titanate AUTHOR (S): Lee, Hyun-Cheol; Jung, Hyun; Oh, Jae-Min; Choy, Jin-Ho CORPORATE SOURCE: National Nanohybrid Materials Laboratory, School of Chemistry and Molecular Engineering, Seoul National University, Seoul, 151-747, S. Korea SOURCE: Bulletin of the Korean Chemical Society (2002), 23(3), 477-480 CODEN: BKCSDE; ISSN: 0253-2964 PUBLISHER: Korean Chemical Society DOCUMENT TYPE: Journal LANGUAGE: English A new microporous TiO2-pillared layered titanate has been successfully prepd. by hybridizing the exfoliated titanate with the anatase TiO2 nano-sol. According to the x-ray diffraction anal. and N2 adsorption-desorption isotherms, the TiO2-pillared layered titanate showed a pillar height of .apprx.2 nm with a high surface area of .apprx.460 m2/g and a pore size of .apprx.0.95 nm, indicating that a microporous pillar structure is formed. Its photocatalytic activity was evaluated by measuring the photodegrdn. rate of 4-chlorophenol during irradn. of catalyst suspensions in an aq. soln. An enhancement in activity of ca. 170% was obtained for TiO2-pillared layered titanate compared to that of the pristine compd. such as layered cesium titanate.

THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS

REFERENCE COUNT:

28

L10 ANSWER 14 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2001:839467 CAPLUS

DOCUMENT NUMBER: 136:38857

TITLE: Functional coatings on basis of inorganic

nanosols Bottcher, H.

AUTHOR (S): CORPORATE SOURCE:

Feinchemie GmbH Sebnitz, Sebnitz, D-01855, Germany Materialwissenschaft und Werkstofftechnik (2001),

32(10), 759-766

CODEN: MATWER; ISSN: 0933-5137

PUBLISHER: Wiley-VCH Verlag GmbH DOCUMENT TYPE: Journal; General Review

LANGUAGE: German

A review. The controlled hydrolysis of silicon or metal alkoxides produces nanoparticulate oxide sols which condense to thin transparent gel films on any substrates after coating and drying (so-called sol-gel process). The co-hydrolysis and co-condensation of different alkoxides (chem. modification) as well as the embedding of different additives (phys. modification) offers almost unlimited possibilities to vary the properties of nanosols and, therefore, also of the resulting coatings, and to adapt them to the purpose intended. By coating of flexible substrates like textiles, papers, or polymer foils it is possible to combine the material protecting functions of the inorg. oxide layer with new functional qualities, e.g. modification of surface energy and charge, alteration of the optical properties, realization of biocompatible and bioactive properties.

REFERENCE COUNT: THERE ARE 66 CITED REFERENCES AVAILABLE FOR THIS 66 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 15 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2001:765446 CAPLUS

DOCUMENT NUMBER:

136:106399

TITLE: Sol-gel processed TiO2-based nano-sized powders for

use in thick-film gas sensors for atmospheric

pollutant monitoring

Traversa, Enrico; Di Vona, Maria Luisa; Licoccia, Silvia; Sacerdoti, Michele; Carotta, Maria Cristina; AUTHOR(S):

Crema, Luigi; Martinelli, Giuliano

CORPORATE SOURCE: Consorzio Interuniversitario Nazionale per la Scienza

e Tecnologia dei Materiali (INSTM) and Dipartimento di Scienze e Tecnologie Chimiche, Universita di Roma "Tor

Vergata", Rome, 00133, Italy

SOURCE: Journal of Sol-Gel Science and Technology (2001),

22(1/2), 167-179

CODEN: JSGTEC; ISSN: 0928-0707 Kluwer Academic Publishers

PUBLISHER: DOCUMENT TYPE: Journal

LANGUAGE: English

Sol-gel routes were used to prep. pure and 5 at.% and 10 at.% Ta- or Nb-dope TiO2 nano-sized powders. The thermal decompn. behavior of the precursors was studied using simultaneous thermogravimetric and DTA (TG/DTA). XRD anal. showed that the powders heated to 400.degree. were cryst. in the anatase TiO2 structure. The pure TiO2 powder heated to 850.degree. showed the rutile structure. The addn. of Ta and Nb inhibited the anatase-to-rutile phase transformation up to 950-1050.degree.. Ta was sol. in the titania lattice up to the concn. of 10 at.%, while the soly. of Nb was 5 at.%. Thick films were fabricated with these powders by screen printing technol. and then fired for 1 h at different temps. in the 650-1050.degree. range. SEM observations showed that the anatase-to-rutile phase transformation induces a grain growth of about one order of magnitude for pure TiO2. The addn. of Ta and Nb is effective to keep the TiO2 grain size at a nanometric level even at 950.degree., though grain growth was obsd. with increasing temp. The gas-sensitive elec. response of the thick films were tested in lab., in environments with CO in dry and wet air. Conductance measurements showed a good gas response only for the nanostructured titania-based films. For field tests, the prototype sensors were placed beside a conventional station for atm. pollutant monitoring. The elec. response of the thick films was compared with the results of the anal. instruments. The same trend was obsd. for both systems, demonstrating the use of gas sensors for this aim.

REFERENCE COUNT: 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 16 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2001:750858 CAPLUS

DOCUMENT NUMBER:

136:9697

TITLE: A new way to prepare nano scale complicated oxide ceramics using an amorphous heteronuclear complex

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AUTHOR (S):
                           Zhu, Y. F.; Yi, T.; Gao, S.; Yan, C. H.; Cao, L. L.
 CORPORATE SOURCE:
                           Department of Chemistry, Tsinghua University, Beijing,
                           100084, Peop. Rep. China
                           Proceedings of the China International Conference on
 SOURCE:
                          High-Performance Ceramics, 1st, Beijing, China, Oct.
                          31-Nov. 3, 1998 (1999), Meeting Date 1998, 162-165. Editor(s): Yan, Dongsheng; Guan, Zhenduo. Tsinghua University Press: Beijing, Peop. Rep. China.
                          CODEN: 69BWTP
 DOCUMENT TYPE:
                          Conference
 LANGUAGE:
                          English
      A new way to prep. nano scale complicated oxide materials using amorphous
      heteronuclear complex as a precursor has been carried out in this paper.
      Gd2CuO4 cuprate oxide was prepd. using this way. The precursor was
      Gd2Cu(DTPA)1.6.6H2O amorphous complex: DTA and TGA indicated that the
      precursor can be decompd. into oxide below 500.degree.. XPS indicated
      that the Gd2CuO4 cuprate oxide were formed after the precursor was
      calcined at 500.degree. for 2 h. XRD also proved the cryst. phase of
      Gd2CuO4 cuprate can be obtained when the calcination temp. increased to
      650.degree.. The cryst. size of Gd2CuO4 increased linearly from 20 nm to
      50 nm when the calcination temp. increases from 500.degree. to 800.degree.
      for 2 h, and the size increases from 10 nm to 30 nm when the calcination
      time increased from 1 to 8 h at 500.degree..
 REFERENCE COUNT:
                                THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS
                          5
                                RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
 L10 ANSWER 17 OF 47 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER:
                          2001:394284 CAPLUS
DOCUMENT NUMBER:
                          135:6781
                          Sol-gel coating on textiles
TITLE:
AUTHOR (S):
                          Bottcher, H.
                          Feinchemie GmbH Sebnitz, Sebnitz, D-01855, Germany
CORPORATE SOURCE:
SOURCE:
                          Textilveredlung (2001), 36(3/4), 16, 19-21
                          CODEN: TXLVAE; ISSN: 0040-5310
PUBLISHER:
                          Verlag Textilveredlung AG
DOCUMENT TYPE:
                          Journal; General Review
LANGUAGE:
                          German
     A review without refs. (list of refs. can be ordered or downloaded) is
     given on prepn. and modification of SiO2-based nano-sols
      and their use for functionalization of textiles: modification of surface
     and optical properties as well as immobilization and controlled release of
     bioactive substances (biocides, fragrances).
L10 ANSWER 18 OF 47 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER:
                         2001:320025 CAPLUS
DOCUMENT NUMBER:
                          134:327989
TITLE:
                         Abrasion-resistant coatings as diffusion barriers
INVENTOR (S):
                         Mennig, Martin; Oliveira, Peter W.; Schmidt, Helmut
PATENT ASSIGNEE(S):
                         Institut Fuer Neue Materialien Gem. G.m.b.H., Germany
SOURCE:
                          PCT Int. Appl., 29 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
     PATENT NO.
                    KIND DATE
                                           APPLICATION NO. DATE
     WO 2001030922
                     A2
                            20010503
                                           WO 2000-EP10589 20001027
     WO 2001030922
                      A3 20020110
        YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
             CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     BR 2000015027
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                            20020716
                                           BR 2000-15027
                                                             20001027
                                           EP 2000-972876 20001027
     EP 1230040
                            20020814
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
    IE, SI, LT, LV, FI, RO, MK, CY, AL
JP 2003512921 T2 20030408 JP 20
                     T2 20030408
                                           JP 2001-533909
                                                            20001027
PRIORITY APPLN. INFO.:
                                        DE 1999-19952040 A 19991028
                                        WO 2000-EP10589 W 20001027
    The title coatings, with high strength in thin layers, comprise base
     layers of polymers which can be cured thermally or photochem. and outer
    layers of compns. contg. nanoscale sols or solid particles which are
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applied on the base layer while the latter is still reactive.

Polycarbonate plates were spin-coated with a 2% iso-PrOH soln. of (EtO)3Si(CH2)3NH2, cured at 130.degree. in dry air, spin-coated with a hydrolyzable epoxysilane, cured at 80.degree. in dry air, spin-coated with a nanosol, and cured at 130.degree. to give a coating with thickness 200-300 nm, good adhesion and abrasion resistance, and moisture diffusion rate .ltoreq.20% lower than that of an uncoated substrate.

L10 ANSWER 19 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:162775 CAPLUS

DOCUMENT NUMBER:

134:213239

TITLE:

In situ XAFS study at the Zr K-edge for SiO2/ZrO2

AUTHOR (S):

CORPORATE SOURCE:

Choy, Jin Ho; Yoon, Joo Byoung; Park, Joo Hyoung School of Chemistry and Molecular Engineering, National Nanohybrid Materials Laboratory, Seoul National University, Seoul, 151-742, S. Korea

SOURCE:

Journal of Synchrotron Radiation (2001), 8(2), 782-784

CODEN: JSYRES; ISSN: 0909-0495

PUBLISHER:

Munksgaard International Publishers Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English

The structural characterization of SiO2/ZrO2 nanosol particles, prepd. by mixing SiO2 sol and aq. soln. of ZrOCl2.8H2O, was carried out by in-situ XAS measurement at the Zr K-edge during condensation reaction. The detailed XANES features at the Zr K-edge of the mixed sol of SiO2/ZrO2 are compared with those of other refs. such as ZrO2, ZrOC12.8H2O, BaZrO3, and ZrSiO4, and it becomes obvious that the Zr4+ ions are stabilized in an octahedral symmetry. Each Zr atom is coordinated with 6 O ones as the first nearest neighbor, where 2 O atoms are from the linkage of (Si-O-Zr) at short distance, and 4 ones are from water mols. at long distance. As the condensation reaction proceeds, it is found that the no. of O atoms due to the formation of (Si-O-Zr) bond at short distance and the second neighbor of silicon atoms increase simultaneously. From the above EXAFS and XANES results, the structural and gelating models could be proposed, which is based on the octahedrally coordinated but distorted Zr species attaching on the SiO2 sol surface.

REFERENCE COUNT:

THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 20 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2001:162757 CAPLUS

14

DOCUMENT NUMBER:

135:83533

TITLE:

Local structure analysis of Ti species stabilized in ion exchangeable layer solids by x-ray absorption

spectroscopy

AUTHOR (S):

Han, Yang Su; Yoon, Sun Mi; Kim, Dong Kuk; Lee, Eun

CORPORATE SOURCE:

SOURCE:

Jung; Choy, Jin Ho; Park, Jung Chul
Department of Chemistry, Kyungpook National
University, Taegu, 702-701, S. Korea
Journal of Synchrotron Radiation (2001), 8(2), 728-730

CODEN: JSYRES; ISSN: 0909-0495

PUBLISHER: Munksgaard International Publishers Ltd. DOCUMENT TYPE: Journal

LANGUAGE: English

Nano-sized TiO2 are incorporated into the interlayer spaces of ion-exchangeable layered perovskites, H1-xCa2-xLaxNb3010 (x = 0.0-0.75), by replacing the interlayer protons with pos. charged TiO2 nanosol particles or basic Ti glycolate complex (titanatrane). Powder x-ray diffraction anal., UV-visible absorption spectroscopy, TGA, and N2 adsorption-desorption isotherm measurements show that quantum sized TiO2 particles are stabilized in between perovskite lattices to form micropores (SBET= 37-110 m2/g). X-ray absorption spectroscopy at the Ti K-edge was used for studying the local environment around Ti atoms constituting the interlayer pillars. According to the XANES spectra, the as-pillared Ti species have the same local environments with those of precursory species, which subsequently converted into TiO2 clusters with rutile and anatase-like local structures when TiO2 nano-sol particles and titanatrane are used as pillaring species, resp. The local environment of TiO2 remains almost const. irresp. of the layer charge d., while the TiO2 pillar content and the microporosity increase with the latter. 8

REFERENCE COUNT:

THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 21 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER:

2000:730908 CAPLUS

DOCUMENT NUMBER: TITLE:

134:12789 Sol-hydrothermal synthesis of SnO2 nano-crystalline

powder

AUTHOR (S):

Lin, Bizhou

CORPORATE SOURCE: College of Chem. Eng., Huaqiao Univ., Quanzhou,

362011, Peop. Rep. China

SOURCE: Huaqiao Daxue Xuebao, Ziran Kexueban (2000), 21(3),

268-270

CODEN: HDZIEF; ISSN: 1000-5013

PUBLISHER: Huagiao Daxue

DOCUMENT TYPE: Journal LANGUAGE: Chinese

Taking SnCl4.cntdot.5H2O as raw material, uniform sized SnO2 nano-cryst. powder with av. grain size of 2.5.apprx.4.0 nm were prepd. by sol-hydrothermal combined method. The nano-cryst. powder were characterized by applying such means as X-ray diffraction, DTA and IR absorption spectrum. The synthesized SnO2 can be used for prepn. of semiconductor sensor for toxic or combustable gases.

L10 ANSWER 22 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2000:24539 CAPLUS

DOCUMENT NUMBER: 132:197830

TITLE: The formation of zirconium hydroxide nanoparticles

from aqueous nitrate solutions

AUTHOR (S): Southon, P. D.; Bartlett, J. R.; Finnie, K. S.;

Woolfrey, J. L.; Ben-Nissan, B.; Kannangara, G. S. K. CORPORATE SOURCE: Department of Chemistry, University of Technology,

Sydney, 2007, Australia

SOURCE: Journal of the Australasian Ceramic Society (1999),

35(1/2), 7-12

CODEN: JAUSEL; ISSN: 1018-6689 PUBLISHER: Australasian Ceramic Society

DOCUMENT TYPE: Journal LANGUAGE: English

Concd. zirconia nano-sols have been prepd. by dissolving Zr oxycarbonate in an acidic soln. of zirconyl nitrate. Hydrolysis and condensation of polynuclear zirconyl cations promotes the formation of nano-sized, polymeric, oxy-hydroxide particles. The evolution of the nanoparticle structure during the formation of the sol has been studied with a range of complementary characterization techniques. Dynamic light scattering indicates that the vast majority of particles are of size 3-6 nm. The Raman spectra of the zirconyl nitrate, and the sol, have been assigned with the assistance of vibrational-modeling software. The characteristic Raman peaks of the polynuclear zirconyl cations at 450 and 575 cm-1 shift to 375 and 535 cm-1, resp., as the particles are formed, indicating the condensation of terminal-hydroxy groups to form hydroxy bridges. 14N NMR and Raman spectroscopy show that all of the unassocd. nitrate groups present in the precursor soln. become weakly assocd. with the surface of the particles,

but are not directly coordinated to the Zr atoms. REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 23 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1999:220251 CAPLUS

DOCUMENT NUMBER: 130:346331

TITLE: Acidic and Hydrophobic Microporous Clays Pillared with

Mixed Metal Oxide Nano-Sols

AUTHOR (S): Han, Yang-Su; Yamanaka, Shoji; Choy, Jin-Ho

Department of Chemistry, Seoul National University, CORPORATE SOURCE:

Seoul, 151-742, S. Korea

SOURCE: Journal of Solid State Chemistry (1999), 144(1), 45-52

CODEN: JSSCBI; ISSN: 0022-4596

PUBLISHER: Academic Press Journal

DOCUMENT TYPE: English

SiO2-metal oxide sol pillared clays were synthesized from montmorillonite by exchanging interlamellar (Na+) ions with SiO2 sol particles modified with polyhydroxy metal cations. Though the SiO2 sol particle itself is neg. charged in the pH range used in present expt., 1.5-2.7, the ion-exchange-type intercalation of the SiO2 sol into montmorillonite was realized by modifying the surface charge of the sol particles from neg. to pos. The pos. charged SiO2 sol particles were prepd. by titrating metal aq. solns. (Mz+ = Fe3+, Al3+, Cr3+, and Zr4+) with NaOH in the presence of SiO2 sol particles, which were easily intercalated in between the silicate layers of clay. On pillaring of oxide sols and subsequent calcining at 400.degree., new porous materials were obtained with high BET surface areas of 320-720 m2/g, pore vols. of 0.24-0.50 mL/g, and basal spacings in the range 40-60 .ANG. Also, their thermal stability could be remarkably improved up to 700.degree. According to the adsorption measurements of N and solvent vapors, the micropores in the samples of size 9-13 .ANG. dominate due to the multilayer stacking of interlayer sol particles in between silicate layers. Temp.-programmed desorption (TPD) spectra of NH3 revealed that the microporous samples are weakly acidic, but with

different strengths, depending on the doped metal species. (c) 1999

Academic Press.

REFERENCE COUNT: THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 24 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1998:750959 CAPLUS DOCUMENT NUMBER: 129:321811

TITLE: Highly porous pillared clay with multistacked

SiO2/TiO2 nanosols

AUTHOR (S): Choy, Jin-Ho; Park, Joo-Hyoung; Yoon, Joo-Byoung

CORPORATE SOURCE: Department of Chemistry, Center for Molecular

Catalysis, College of Natural Sciences, Seoul National

University, Seoul, 151-742, S. Korea

SOURCE: Bulletin of the Korean Chemical Society (1998),

19(11), 1185-1188

CODEN: BKCSDE; ISSN: 0253-2964

PUBLISHER: Korean Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

Layered nanocomposite, SiO2/TiO2 sol pillared clay, has been prepd. by the ion exchange reaction of Na+ ion in montmorillonite with pos. charged mixed SiO2/TiO2 sol. The nanosized sol particles were synthesized by mixing SiO2 sol soln. with TiO2 one, which is obtained by acidic hydrolysis of TEOS and TiCl4, resp. From powder XRD, the basal spacing (d001) of the sample calcined at 400 .degree.C was found to be ca. 60 .ANG., due to the multistacking of nanosized SiO2 and TiO2 sol particles, which was confirmed by the pore size anal. from 129Xe NMR and micropore anal. calcd. from nitrogen adsorption. The BET sp. surface area shows the value of 684 m2g-1 (Langmuir 1115 m2g-1), which is the highest among various pillared clays ever reported previously, and the total porosity is found to be 0.51 mlg-1, and the pores are mainly composed of micropore with a size of ca. 11.8 .ANG.. This result agrees with the adsorption capacity obtained from water adsorption. According to diffuse reflectance UV-visible spectroscopy, it is found that the TiO2 particles stabilized in the interlayer space of montmorillonite are quantum-sized of ca. 20 .ANG..

REFERENCE COUNT: 22 THERE ARE 22 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 25 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1998:589936 CAPLUS

DOCUMENT NUMBER: 129:279493

TITLE: Sol-Gel Preparation of AuCu and Au4Cu Nanocluster

Alloys in Silica Thin Films

AUTHOR (S): Gwak, Ji-Hye; Kim, Sung-Jin; Lee, Minyung CORPORATE SOURCE:

Department of Chemistry, Ewha Womans University,

Seoul, 120-750, S. Korea

Journal of Physical Chemistry B (1998), 102(40), SOURCE:

7699-7704

CODEN: JPCBFK; ISSN: 1089-5647

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

Using sol-gel processing and high-temp. annealing, metal alloy nanoclusters of AuCu and Au4Cu in SiO2 thin films were synthesized. The crystallog. data show that the lattice parameters of those nanocluster alloys, both showing the fcc. structure, are very close to the bulk values. The nanocrystals obtained by annealing at 900.degree. were nearly spherical in shape, having the AuCu size of .apprx.23 nm in radius and the Au4Cu size of .apprx.35 nm. The peaks of the optical spectra of the samples annealed at 700-900.degree. are located at 555 nm for AuCu and at 562 nm for Au4Cu.

REFERENCE COUNT: 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 26 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1998:587130 CAPLUS

DOCUMENT NUMBER: 129:222968

TITLE: Optical switching and limiting: new applications for

silver halide technology?

Sahyun, M. R. V.; Hill, Susan E.; Serpone, N.; Danesh, Reza; Sharma, Devendra K. AUTHOR (S):

CORPORATE SOURCE: Department Chemistry, University Wisconsin, Eau

Claire, WI, USA

SOURCE: IS&T's Annual Conference (1997), 50th, 19-20

CODEN: ISACFN

PUBLISHER: Society for Imaging Science and Technology

DOCUMENT TYPE: Journal LANGUAGE: English

The exptl. result already published imply a biphotonic mechanism of

spectral sensitization. Anal. of the optical limiting mechanism suggests that iodide doping, which is adventitious in the present case, is crit. to the obsd. photophys. behavior of the nanosols. The reaction

sequence of rapid optical switching by Ag halide was presented.

REFERENCE COUNT: THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS 11 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 27 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1998:447109 CAPLUS

DOCUMENT NUMBER:

129:127565

TITLE:

Multilayered SiO2/TiO2 Nanosol Particles in

AUTHOR(S):

PUBLISHER:

Two-Dimensional Aluminosilicate Catalyst-Support Choy, Jin-Ho; Park, Joo-Hyoung; Yoon, Joo-Byoung

CORPORATE SOURCE: Department of Chemistry Center for Molecular Catalysis College of Natural Sciences, Seoul National

University, Seoul, 151-742, S. Korea

SOURCE:

Journal of Physical Chemistry B (1998), 102(31),

5991-5995

CODEN: JPCBFK; ISSN: 1089-5647 American Chemical Society

DOCUMENT TYPE:

Journal

LANGUAGE:

English

A new layered nanocomposite, which is one-to-one interstratified with a montmorillonite layer and a mixed SiO2/TiO2 sol particle one, has been prepd. by ion exchange reaction of the Na+ ion in montmorillonite with the pos. charged SiO2/TiO2 sol particles. The ion exchange reaction was performed at three different temps. of 45, 60, and 75 .degree.C by mixing an aq. suspension of 1 wt % Na+ montmorillonite with SiO2/TiO2 sol soln. where the molar ratio of Si/Ti was selected as 20/2. According to the powder X-ray diffraction anal., the basal spacings of layered nanocomposites calcined at 400 .degree.C were found to increase from 35.4 ANG., to 47.3 ANG., and to 60.0 ANG. as the ion exchange reaction temp. was raised from 40 .degree.C, to 60 .degree.C, and to 75 .degree.C. Their BET and Langmuir sp. surface areas and porosities, estd. from nitrogen adsorption-desorption isotherms, become larger with the increment of basal spacing, and the highest BET sp. surface area and the largest porosity are found to be 683 m2/g and 0.50 mL/g, resp. Despite the large increment of the basal spacing, the porous properties such as sp. surface areas, porosities, and pore sizes, those which are calcd. from t-plots and chem. shift of 129Xe NMR, resp., are detd. to be almost const. From the UV/vis spectra, the blue shift of the absorption edge was obsd., indicating that the TiO2 sol particles in the interlayer are quantum sized. It is therefore proposed that the products are intercalation-type nanocomposites with the multistacked structure of the SiO2/TiO2 nanoparticles in the interlayer space of montmorillonite.

REFERENCE COUNT:

THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS 30 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 28 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: DOCUMENT NUMBER:

1997:711691 CAPLUS

127:286203

TITLE:

Sol-Gel Template Synthesis of Semiconductor Oxide

Micro- and Nanostructures

AUTHOR (S):

Lakshmi, Brinda B.; Patrissi, Charles J.; Martin,

Charles R.

CORPORATE SOURCE:

Department of Chemistry, Colorado State University, Fort Collins, CO, 80523, USA

PUBLISHER: DOCUMENT TYPE: Chemistry of Materials (1997), 9(11), 2544-2550 CODEN: CMATEX; ISSN: 0897-4756

American Chemical Society Journal; General Review

SOURCE:

English

The template method for synthesizing nanostructures involves the synthesis of the desired material within the pores of a nanoporous membrane or other solid. The authors' work has involved using porous alumina and polymeric filter membranes as the templates. Fibrils or tubules of the desired material are formed within each pore of the template membrane. A no. of synthetic methods were used to synthesize these nanostructures. This paper reviews with many refs. sol-gel template synthesis: the use of sol-gel chem. to synthesize semiconductor oxide micro- and nanostructures within the pores of micro- and nanoporous membranes. For example, TiO2 nanotubules and nanofibers of the anatase form were synthesized. The high surface area offered by these TiO2 nanostructures was used for photodecompn. of salicylic acid in sunlight. Enzyme immobilization by stannous bridges inside the TiO2 tubes also was studied. V2O5 fibrous electrode materials were prepd. by this method and Li intercalation electrochem. is reported here. Other semiconductor oxides such as MnO2, Co3O4, ZnO, WO3, and SiO2 also were prepd.

ACCESSION NUMBER: 1997:298944 CAPLUS

DOCUMENT NUMBER: 127:22124

TITLE: Preparation and optical properties of TiO2

nanocrystalline particles dispersed in SiO2

nano-composites

AUTHOR (S): Zhou, Q. F.; Zhang, Q. Q.; Zhang, J. X.; Zhang, L. Y.;

Yao, X.

Department of Physics, Zhongshan University, Canton, CORPORATE SOURCE:

510275, Peop. Rep. China

SOURCE: Materials Letters (1997), 31(1,2), 39-42

CODEN: MLETDJ; ISSN: 0167-577X

PUBLISHER: Elsevier DOCUMENT TYPE: Journal LANGUAGE: English

Nano-composites of TiO2 nanocryst. particles dispersed in SiO2 have been successfully prepd. by the two-step hydrolysis of Si(OC2H5)4 derived by the sol-gel technique using TiO2 ultrafine powder as a active component. The influence of processing factors has been investigated; a two-step rapid gelation method in sol-gel processing has been developed. The optical absorption spectrum and the third-order nonlinear susceptibility .chi.(3) of the nano-composites were measured, the highest .chi.(3) of the sample was 4.1.times.10-11 esu measured by the DFWM method.

L10 ANSWER 30 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: CAPLUS

1997:236959 DOCUMENT NUMBER: 126:320003

TITLE: Nanocomposites Fe2O3/SiO2 - Preparation by sol-gel

method and physical properties

AUTHOR (S): Niznansky, D.; Viart, N.; Rehspringer, J.L.

Institute of Inorganic Chemistry ASCR, Rez, 250 68, CORPORATE SOURCE:

Czech Rep.

SOURCE: Journal of Sol-Gel Science and Technology (1997),

8(1/2/3), 615-618

CODEN: JSGTEC; ISSN: 0928-0707

PUBLISHER: Kluwer DOCUMENT TYPE: Journal LANGUAGE: English

Magnetic nanocomposites .gamma.-Fe2O3/silica were prepd. by a one-step sol-gel method. The sol was prepd. by TEOS (tetra-Et orthosilicate) acid hydrolysis in the presence of an iron salt sol. in methanol. After gelation and drying, the transparent samples were characterized after treatment at different temps. The particle size, obsd. by HR-TEM, was in the range of 2-10 nm and depended on Fe-concn. and heating temp. Magnetic measurements showed either a ferromagnetic or a superparamagnetic behavior and could be explained by the particle size. The dependence of the magnetic behavior on the particle size was also studied by Mossbauer spectroscopy. The samples in which the Fe203 particle size was approx. 10 nm showed magnetic splitting (sextet) at room temp., while smaller particles (2-3 nm) showed this splitting only at the temp. of liq. helium. The optomagnetic properties of the samples were also measured (Kerr effect).

L10 ANSWER 31 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1997:134897 CAPLUS

DOCUMENT NUMBER: 126:160941

TITLE:

Sol-Gel Template Synthesis of Semiconductor

Nanostructures

AUTHOR (S): Lakshmi, Brinda B.; Dorhout, Peter K.; Martin, Charles

CORPORATE SOURCE: Department of Chemistry, Colorado State University, Fort Collins, CO, 80523, USA

SOURCE: Chemistry of Materials (1997), 9(3), 857-862

CODEN: CMATEX; ISSN: 0897-4756

PUBLISHER: American Chemical Society DOCUMENT TYPE: Journal

LANGUAGE: English

The template method for prepg. nanostructures entails synthesis of the desired material within the pores of a nanoporous membrane or other solid. A nanofibril or tubule of the desired material is obtained within each pore. Methods used previously to deposit materials within the pores of such membranes include electrochem. and electroless deposition and in situ polymn. This paper describes the first use of sol-gel chem. to prep. semiconductor nanofibrils and tubules within the pores of an alumina template membrane. TiO2, WO3, and ZnO nanostructures have been prepd. TiO2 nanofibrils with diams. of 22 nm were found to be single crystals of anatase with the c-axis oriented along the fibril axis. Bundles of these fibrils were also found to be single cryst., suggesting that the individual fibrils are arranged in a highly organized fashion within the bundle. Finally, 200 nm diam. TiO2 fibrils were used as photocatalysts for the decompn. of salicylic acid.

L10 ANSWER 32 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1996:293907 CAPLUS

DOCUMENT NUMBER:

124:356021

TITLE: Optical limiting characteristics and mechanism of

silver bromide nanosols

AUTHOR (S): Sahyun, M. R. V.; Hill, Susan E.; Serpone, N.; Danesh,

Reza; Sharma, Devendra K.

CORPORATE SOURCE: Dry Imaging Technology Center, 3M 3M Center, St. Paul,

MN, 55144, USA

SOURCE: Journal of Applied Physics (1996), 79(10), 8030-8037

CODEN: JAPIAU; ISSN: 0021-8979

PUBLISHER: American Institute of Physics

DOCUMENT TYPE: Journal LANGUAGE: English

Optical limiting behavior has been obsd. in nanosols comprising .apprx.60 .ANG. particles of silver bromide. Switching times, .tau., are consistently in the ns regime, and values of I1/2, the pulse laser exposure which elicits a 50% decrease in transmittance of the medium, can be less than 20 mJ/cm2. Nonlinearity of response, I1/2, and response time all increase with decreasing pAg of the nanosols. The materials can be cycled apparently indefinitely, provided they are subjected only to very short (sub-ns) laser pulses. Optical switching can be spectrally sensitized; comparable response parameters are obtained under these conditions. The nonlinear character of the response is, however, very different: a true response threshold, IO = .apprx. 12 mJ/cm2, is obsd. Anal. of the optical limiting mechanism suggests that iodide doping, which is adventitious in the present case, is crit. to the obsd. photophys. behavior of the nanosols.

L10 ANSWER 33 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER:

DOCUMENT NUMBER:

1996:139722 CAPLUS 124:209625

TITLE:

SOURCE:

AUTHOR (S):

Fabrication, characterization and application of Ni/SiO2 nanocomposite materials prepared by sol-gel Monaci, R.; Musinu, A.; Piccaluga, G.; Pinna, G.

CORPORATE SOURCE:

Dipartimento di Scienze Chimiche, Cagliari, I-09124, Italy

Materials Science Forum (1995), 195 (Nanophase Materials), 1-6

CODEN: MSFOEP; ISSN: 0255-5476

PUBLISHER: Trans Tech

DOCUMENT TYPE: Journal LANGUAGE · English

Composites of nanoparticles of nickel dispersed in amorphous silica have been prepd. by classical sol-gel processing. Mild conditions have been used whenever possible to prevent the formation of large particles of nickel; times and temps. of the redn. step were varied in the treatment of samples with 7% nickel to verify their influence on the particle size. The samples with Ni particles of about 5 nm showed good catalytic activity in a std. hydrogenation reaction.

L10 ANSWER 34 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER:

DOCUMENT NUMBER:

1996:17143 CAPLUS 124:131239

TITLE:

SOURCE:

Mechanisms of spectral sensitization: Role of

sensitizing dye complexation

AUTHOR (S): CORPORATE SOURCE: Sahyun, M. R. V.; Sharma, D. K.; Serpone, N. Information, Imaging and Electronics Sector

Laboratories 3M, St. Paul, MN, USA

IS&T's Annual Conference: Imaging on the Information Superhighway, Final Program and Advance Printing of Papers, 48th, Washington, D.C., May 7-11, 1995 (1995), 219. IS&T--The Society for Imaging Science and

Technology: Springfield, Va.

CODEN: 62BXAS

DOCUMENT TYPE:

Conference LANGUAGE: English

We have obsd. concurrent processes of photoexcited dye deactivation and Ag(0) cluster formation in real time under conditions of laser flash photolysis of a merocyanine dye adsorbed to an AgBr nanosol. To our knowledge this is the first time such a comprehensive view of the process of spectral sensitization has been obtained in an exptl. system.

L10 ANSWER 35 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER:

1995:946448 CAPLUS

DOCUMENT NUMBER:

124:71337

TITLE:

Mechanisms of spectral sensitization of silver halides: role of sensitizing dye complexation Sahyun, M. R. V.; Sharma, D. K.; Serpone, N.

AUTHOR (S):

CORPORATE SOURCE: Dry Imaging Technol. Cent., St. Paul, MN, 55144, USA SOURCE:

Journal of Imaging Science and Technology (1995).

39(5), 377-85

CODEN: JIMTE6; ISSN: 1062-3701

PUBLISHER: IS&T--The Society for Imaging Science and Technology

DOCUMENT TYPE: Journal LANGUAGE: English

We have obsd. concurrent processes of photoexcited dye deactivation and silver(0) cluster formation in real time under conditions of laser flash photolysis of a merocyanine dye adsorbed to an AgBr nanosol. To our knowledge this is the first time such a comprehensive view of the process of spectral sensitization has been obtained exptl. Spectral sensitization of AgBr is apparently biphotonic under our conditions; this result, along with obsd. kinetics of dye ground state re-population and of silver(0) cluster growth, is consistent with Mitchell's mechanism of spectral sensitization, but not with single-electron transfer or radical pair mechanisms. Control expts. revealed a modicum of photolytic reactivity for undyed nanosol with 2.35 eV photons and suggested operation of an Auger mechanism of photoelectron generation. In this case amplified stimulated emission was obsd. from photogenerated silver clusters, AgnO, (or a byproduct) at photon energies comparable to those that produce the Herschel effect in conventional photog. We speculatively est. nuclearity of the silver clusters produced under conditions of our expts. as n = ca. 12.

L10 ANSWER 36 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1995:222141 CAPLUS

DOCUMENT NUMBER: 122:15318

TITLE: Sol-gel processing of inorganic membranes

AUTHOR (S): Guizard, C.; Mouchet, C.; Vacassy, R.; Julbe, A.;

Larbot, A.

CORPORATE SOURCE: Laboratoire des Materiaux et des Procedes

Membranaires, Ecole Nationale Superieure de Chimie.

Montpellier, 34053, Fr.

SOURCE: Journal of Sol-Gel Science and Technology (1994), 2(1/2/3), 483-7

CODEN: JSGTEC; ISSN: 0928-0707 PUBLISHER: Kluwer

DOCUMENT TYPE: Journal LANGUAGE: English

Basic principles involved in sol-gel processing of ceramic membranes are described. This process has been applied to ceramic ultrafiltration membranes and is now investigated to prep. ceramic nanofilters. Special emphasis is put on new developments concerning microporous zirconia membranes obtained by the polymeric route. A zirconium alkoxide precursor modified with an acetylacetone ligand has been used to control particle growth in the sols and pore size distribution in the membranes. N2 adsorption and x-ray diffraction anal. have been performed on membrane materials showing the influence of process parameters (molar ratio .gamma. = acacH/Zr and sintering temp. T) on membrane structural evolution.

L10 ANSWER 37 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1993:676990 CAPLUS DOCUMENT NUMBER: 119:276990

TITLE: Creating better nanocomposites

AUTHOR (S): Lukehart, Charles M.; Carpenter, Joseph P.; Milne,

Stephen B.; Burnam, Kimberly J.

CORPORATE SOURCE: Dep. Chem., Vanderbilt Univ., Nashville, TN, 37235,

SOURCE: CHEMTECH (1993), 23(8), 29-34 CODEN: CHTEDD; ISSN: 0009-2703

DOCUMENT TYPE: Journal; General Review

LANGUAGE: English

A review, with 13 refs., of the incorporation of metal complexes in silica xerogels to prep. highly dispersed nanoclusters of metals or semiconductors supported on SiO2. Discussions include the Co-carbido system, Ag-thiolate complex system, organogermanism compd. system, and optical studies of the Ge nanocomposites.

L10 ANSWER 38 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1993:503346 CAPLUS

DOCUMENT NUMBER: 119:103346

TITLE:

Dihydropyridine derivative-containing nanosol

as oral delayed-release dosage form

INVENTOR(S): Wunderlich, Jens Christian; Schick, Ursula; Werry,

Juergen; Freidenreich, Juergen Alfatec-Pharma GmbH, Germany

Ger. Offen., 8 pp. SOURCE:

CODEN: GWXXBX

DOCUMENT TYPE: Patent

PATENT ASSIGNEE(S):

LANGUAGE: German FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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PATENT NO.
                   KIND DATE
                                         APPLICATION NO. DATE
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    DE 4140194
                     A1
                          19930609
                                         DE 1991-4140194 19911205
                     C2 19980219
    DE 4140194
    WO 9310770
                     A1
                          19930610
                                        WO 1992-DE1014 19921204
        W: AU, CA, JP, US
        RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
                  A1 19930628
A1 19940921
    AU 9230806
                                        AU 1992-30806
                                                        19921204
    EP 615446
                                        EP 1992-924551
                                                        19921204
    EP 615446
                     B1
                         19951115
       R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, NL, PT, SE
              E 19951215
    AT 130190
                                      AT 1992-924551 19921204
    ES 2082524
                                        ES 1992-924551
                     T3
                          19960316
                                                        19921204
PRIORITY APPLN. INFO.:
                                     DE 1991-4140194
                                                        19911205
                                     WO 1992-DE1014
                                                        19921204
```

A dihydropyridine deriv. for treatment of cardiovascular disease is incorporated as the inner phase in a nanosol having an outer phase of gelatin or a gelatin fraction or deriv. bearing an elec. charge opposite to that of the drug so that the sol as a whole is elec. neutral. This prepn. shows improved bioavailability, resorption, and biocompatibility. Thus, nifedipine (30 g in 0.5 L iso-PrOH) was added to a soln. of 600 g of a 6% soln. of type B gelatin (pI 4.7) at 60.degree. and pH 5.5 to form a nanosol, which was spray-dried and pressed into tablets each contg. 20 mg nifedipine.

L10 ANSWER 39 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1993:503344 CAPLUS

DOCUMENT NUMBER:

119:103344

TITLE:

INVENTOR(S):

Delayed-release form of a 3-indoleacetic acid

derivative-containing drug preparation Wunderlich, Jens Christian; Schick, Ursula;

PATENT ASSIGNEE(S):

Freidenreich, Juergen; Werry, Juergen Alfatec-Pharma GmbH, Germany

SOURCE:

Ger. Offen., 7 pp.

CODEN: GWXXBX

Patent

DOCUMENT TYPE: LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 4140191	A1	19930609	DE 1991-4140191	19911205
DE 4140191	C2	19980219		
WO 9310769	A1	19930610	WO 1992-DE1013	19921204
W: AU, CA,	JP, US			
RW: AT, BE,	CH, DE	, DK, ES,	FR, GB, GR, IE, IT, LU,	MC, NL, PT, SE
AU 9230805	A1	19930628	AU 1992-30805	19921204
PRIORITY APPLN. INFO	.:		DE 1991-4140191	19911205

WO 1992-DE1013 19921204 An antirheumatic or anti-inflammatory 3-indoleacetic acid deriv. (e.g. indomethacin, acemethacin) is incorporated as the inner phase in a nanosol having an outer phase of gelatin, a gelatin deriv., or a gelatin fraction bearing an elec. charge opposite to that of the drug so that the sol as a whole is elec. neutral. The 3-indoleacetic acid deriv. is essentially completely resorbed from this prepn. in all portions of the digestive tract independently of the physiol. conditions in different portions of the tract or of the phys.-chem. properties of the compd. Thus, 100 g indomethacin was dissolved in a soln. of 600 g type B gelatin (pI 5.2) in 10 L distd. water at 55.degree. and pH 7-8. Addn. of HCl to pH 3.1 produced a nanosol, which was spray-dried and pressed into tablets each contg. 75 mg indomethacin.

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L10 ANSWER 40 OF 47 CAPLUS COPYRIGHT 2003 ACS
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ACCESSION NUMBER: DOCUMENT NUMBER:

1993:503342 CAPLUS

119:103342

TITLE:

Pharmaceutical preparation containing a 2-arylpropionic acid derivative in nanosol

INVENTOR (S):

Wunderlich, Jens Christian; Schick, Ursula;

Freidenreich, Juergen; Werry, Juergen; Lukas, Helmut;

Schuster, Otto

PATENT ASSIGNEE(S):

Alfatec-Pharma GmbH, Germany; PAZ Arzneimittelentwicklungsgesllschaft mbH

SOURCE:

Ger. Offen., 7 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent LANGUAGE: German FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

```
PATENT NO. KIND DATE
      PATENT NO.
                                                     APPLICATION NO. DATE
     DE 4140185 A1 19930609
DE 4140185 C2 19960201
WO 9310761 A1 19930610
                                                     DE 1991-4140185 19911205
                                                     WO 1992-DE1012 19921204
          W: AU, CA, JP, US
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
      AU 9230804 A1 19930628
AU 669500 B2 19960613
                                                    AU 1992-30804
                                                                          19921204
     AU 669500 B2 19960013
EP 615440 A1 19940921
EP 615440 B1 19950913
                                                     EP 1992-924549
                                                                         19921204
          \mbox{R:} \quad \mbox{AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, NL, PT, SE} \\
      ES 2080527 T3 19960201
CA 2125283 C 19980922
US 5560924 A 19961001
                                                    ES 1992-924549 19921204
                                                     CA 1992-2125283 19921204
                                                     US 1994-244688 19940929
                                  19961001
PRIORITY APPLN. INFO.:
                                                 DE 1991-4140185
                                                                         19911205
                                                 WO 1992-DE1012
                                                                         19921204
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An antipyretic, analgesic, or anti-inflammatory 2-arylpropionic acid deriv. (e.g. ibuprofen, ketoprofen) is incorporated as the inner phase in a nanosol having an outer phase of gelatin, a gelatin deriv., or a collagen hydrolyzate bearing an elec. charge opposite to that of the drug so that the sol as a whole is elec. neutral. The drug is stabilized by the gelatin and is rapidly and completely resorbed in the stomach. Thus, 300 g racemic ibuprofen was dissolved in 800 g 10% NaOH soln. and added to 10 kg of a 6% soln. of gelatin type B (pI 4.9) at 40.degree. to provide a clear soln. This soln. was adjusted to pH 3.0 with HCl to form a nanosol, which was spray dried, mixed with excipients, and pressed into tablets each contg. 200 mg ibuprofen.

L10 ANSWER 41 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER:

1993:480243 CAPLUS

DOCUMENT NUMBER:

119:80243

TITLE:

Rapid-release flurbiprofen nanosol.

INVENTOR (S):

Wunderlich, Jens Christian; Lukas, Helmut; Schuster,

Otto; Schick, Ursula

PATENT ASSIGNEE(S):

Alfatec-Pharma GmbH, Germany; PAZ Arzneimittelentwicklungsgesellschaft mbH

SOURCE: Ger. Offen., 6 pp.

CODEN: GWXXBX

DOCUMENT TYPE:

Patent German

LANGUAGE: FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND DAT	E A	PPLICATION NO.	DATE
DE 4140184	A1 199	30609 DI	F 1991_4146194	10011205
DE 4140184	C2 1999	51221	P 1001-4140104	19911205
WO 9310766	A1 199	30610 W	O 1992-DE1008	19921204
W: AU, CA,	JP, US			
RW: AT, BE,	CH, DE, DK,	ES, FR, GB,	GR, IE, IT, LU	, MC, NL, PT, SE
AU 9229409	A1 1993	30628 At	J 1992-29409	19921204
EP 615443	A1 1994	10921 EI	P 1992-923694	19921204
EP 615443	B1 1996	0814		
R: AT, BE,	CH, DE, DK,	ES, FR, GB,	GR, IT, LI, NL,	, PT, SE
AT 141160	E 1996	0815 AT	Г 1992-923694	19921204
ES 2090703	T3 1996	1016 ES	5 1992-923694	19921204
US 5556638	A 1996	ເດງ17 ປຽ	3 1994-244614	19940916
PRIORITY APPLN. INFO	.:		991-4140184	
			992-DE1008	

A rapid-release anti-inflammatory formulation comprises a flurbiprofen inner phase (10-800 nm particle size) and a gelation or collagen hydrolyzate outer phase. The two phases have opposite isoionic elec charges. A soln. of 600 g gelatin B in 10 L water was blended with a soln. of 200 g R-flurbiprofen in 700 g 10% NaOH, followed by pH adjustment to 3.2 (HCl) to give a nanosol, which was spray-dried and tabletted.

L10 ANSWER 42 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1993:480242 CAPLUS

DOCUMENT NUMBER:

119:80242

TITLE:

Rapid-release ibuprofen nanosol Wunderlich, Jens Christian; Lukas, Helmut; Schuster,

Otto; Schick, Ursula

PATENT ASSIGNEE(S):

INVENTOR (S):

Alfatec-Pharma GmbH, Germany; PAZ

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Arzneimittelentwicklungsgesellschaft GmbH
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SOURCE: Ger. Offen., 9 pp.

CODEN: GWXXBX

DOCUMENT TYPE: LANGUAGE:

Patent German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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PATENT NO.
                     KIND DATE
                                          APPLICATION NO. DATE
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     DE 4140179
                     A1
                           19930609
                                          DE 1991-4140179 19911205
    DE 4140179
                      C2
                           19951221
     WO 9310762
                           19930610
                      A1
                                          WO 1992-DE1016
                                                           19921204
        W: AU, CA, JP, US
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
    AU 9230808
                    A1 19930628
                                          AU 1992-30808
                                                           19921204
    AU 670048
                      B2
                           19960704
    EP 615441
                     Al 19940921
                                          EP 1992-924553
    EP 615441
                      B1
                           19960626
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, NL, PT, SE
    AT 139690
               E 19960715
                                         AT 1992-924553
                                                          19921204
    ES 2090712
                      Т3
                           19961016
                                          ES 1992-924553
                                                          19921204
    CA 2125281
                      C
                           19980922
                                          CA 1992-2125281 19921204
    US 6066332
                                          US 1995-244690
                      Α
                           20000523
                                                          19951201
PRIORITY APPLN. INFO.:
                                       DE 1991-4140179
                                                          19911205
                                       WO 1992-DE1016
                                                          19921204
```

A rapid-release anti-inflammatory formulation comprises an ibuprofen inner phase (10-800 nm particle size) and a gelatin or collagen hydrolyzate outer phase. The 2 phases have opposite isoionic elec. charges. A soln. of 600 g gelatin B in 10L water was blended with a soln. of 300g S-ibuprofen in 0.8 L 10% NaOH, followed by pH adjustment to 3 (HCl) to give a nanosol, which was spray-dried and tabletted.

L10 ANSWER 43 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER:

DOCUMENT NUMBER:

1993:456219 CAPLUS 119:56219

TITLE:

Slow-release flurbiprofen nanosol

INVENTOR (S):

Wunderlich, Jens Christian; Schuster, Otto; Lukas,

Helmut; Schick, Ursula

PATENT ASSIGNEE(S):

Alfatec-Pharma GmbH, Germany; Paz Arzneimittelentwicklungsgesellschaft MbH

SOURCE: PCT Int. Appl., 45 pp.

CODEN: PIXXD2

formulation

DOCUMENT TYPE:

Patent German

LANGUAGE: FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE
*** *****			
WO 9310771	Al 19930610	WO 1992-DE1015	19921204
W: AU, CA,	JP, US		
RW: AT, BE,	CH, DE, DK, ES,	FR, GB, GR, IE, IT, LU	, MC, NL, PT, SE
DE 4140183	Al 19930617	DE 1991-4140183	19911205
DE 4140183	C2 19951221		17711103
AU 9230807	Al 19930628	AU 1992-30807	19921204
EP 615447	A1 19940921	EP 1992-924552	19921204
EP 615447	B1 19960306		13321204
R: AT, BE,	CH, DE, DK, ES,	FR, GB, GR, IT, LI, NL	. PT. SE
AT 134876	E 19960315	AT 1992-924552	
ES 2085657	T3 19960601		
PRIORITY APPLN. INFO.		DE 1991-4140183	
2.1.2	• •		
		WO 1992-DE1015	19921204

19921204 Flurbiprofen (racemate, pseudoracemate or enantiomers) is formulated as a nanosol, for sustained release. A soln. of 600 g gelatin type B in 10 L water was treated at 50.degree., with a soln. of 200 g S-flurbiprofen in 350 g 10% NaOH, followed by pH adjustment to 3.2 (HCl) to give a nanosol, which was spray-dried.

L10 ANSWER 44 OF 47 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER:

1993:456216 CAPLUS

DOCUMENT NUMBER:

119:56216

TITLE:

Slow-release ibuprofen nanosol formulations

INVENTOR (S):

Wunderlich, Jens Christian; Schuster, Otto; Lukas,

Helmut; Schick, Ursula

PATENT ASSIGNEE(S):

Alfatec-Pharma GmbH, Germany; Paz

SOURCE:

Arzneimittelentwicklungsgesselschaft mbH PCT Int. Appl., 49 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: German FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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PATENT NO.
                      KIND DATE
                                               APPLICATION NO. DATE
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                       Al
     WO 9310760
         9310760

W: AU, CA, JP, US

RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE

140172

A1 19930617

DE 1991-4140172 19911205
                              19930610
                                               WO 1992-DE1007 19921204
     DE 4140172 A1 19930617
     DE 4140172
     AU 9230800 A1 19930628
EP 615439 A1 19940921
EP 615439 B1 19951108
                                               AU 1992-30800
                                                                  19921204
                                               EP 1992-924545 19921204
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, NL, PT, SE
     AT 129891 E 19951115
ES 2082523 T3 19960316
                                          AT 1992-924545 19921204
                                               ES 1992-924545
                                                                  19921204
PRIORITY APPLN. INFO.:
                                           DE 1991-4140172
                                                                  19911205
                                            WO 1992-DE1007
                                                                  19921204
```

Ibuprofen (racemate, pseudoracemate, enantiomers) is formulated as a nanosol, for sustained release. A soln. of 300 g gelatin type B in 5 L water was treated with a soln. of 200 g S-ibuprofen in 0.6 L 10% NaOH, followed by pH adjustment to 3 (HCl) to give a nanosol, which was spray-dried.

L10 ANSWER 45 OF 47 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1993:456196 CAPLUS

DOCUMENT NUMBER: 119:56196

TITLE: Rapid-release 3-indolacetic acid derivative

nanosol drug

INVENTOR(S): Wunderlich, Jens Christian; Schick, Ursula; Werry,

Juergen; Freidenreich, Juergen Alfatec-Pharma GmbH, Germany

PATENT ASSIGNEE(S): SOURCE:

Ger. Offen., 6 pp.

DOCUMENT TYPE:

CODEN: GWXXBX Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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PATENT NO. KIND DATE
                                            APPLICATION NO. DATE
                             -----
     DE 4140178 Al 19930609
                                             DE 1991-4140178 19911205
     DE 4140178
                        C2
                             19980219
                      A1 19930610
      WO 9310767
                                             WO 1992-DE1009 19921204
         W: AU, CA, JP, US
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
      WO 9310768
                       A1 19930610
                                           WO 1992-DE1010 19921204
         W: AU, CA, JP, US
         RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
                       A1 19930628
B2 19960919
     AU 9230801
                                            AU 1992-30801 19921204
     AU 671964
     AU 9230802
                      A1 19930628
B2 19960919
                                             AU 1992-30802
                                                               19921204
     AU 671965
                      A1 19940921
     EP 615444
                                             EP 1992-924546 19921204
         615444 B1 19960306
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, NL, PT, SE
     EP 615444
     EP 615445 Al 19940921
                                            EP 1992-924547 19921204
     EP 615445
                       B1
                            19960515
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, NL, PT, SE
                E 19960315 AT 1992-924546 19921204
T3 19960601 ES 1992-924546 19921204
     AT 134875
                                            AT 1992-924546 19921204
ES 1992-924546 19921204
19921204
     ES 2085656
     AT 137962
                       E 19960615
T3 19960716
                       Е
     ES 2087565
     US 5614219
                                             ES 1992-924547 19921204
                       A
                            19970325
                                            US 1994-244691
                                                              19940913
     US 5932245
                       Α
                            19990803
                                            US 1994-244615
                                                              19941025
PRIORITY APPLN. INFO.:
                                         DE 1991-4140177 A 19911205
DE 1991-4140178 A 19911205
                                         DE 1991-4140186 A 19911205
                                         DE 1991-4140195 A 19911205
                                         US 1992-876867 A 19920430
                                         WO 1992-DE1009 A 19921204
WO 1992-DE1010 A 19921204
                                                           A 19921204
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A rapid-release anti-inflammatory nanosol formulation comprises an inner phase made of a title drug (indomethacin or acemethacin), having a 10-600 nm particle size, and a gelatin or collagen hydrolyzate outer phase. The 2 phase, have opposite isoionic charges. A suspension (pH 7.6) of 100 g indomethacin in a soln. of 600 g gelatin in 10L water was adjusted to pH 3.1, followed by spray-drying and tabletting, to give a

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L10 ANSWER 46 OF 47 CAPLUS COPYRIGHT 2003 ACS
  ACCESSION NUMBER:
                            1993:456195 CAPLUS
  DOCUMENT NUMBER:
                            119:56195
  TITLE:
                            Rapid-release glibenclamide nanosol
  INVENTOR (S):
                            Wunderlich, Jens Christian; Schick, Ursula;
                            Freidenreich, Juergen; Werry, Juergen
  PATENT ASSIGNEE(S):
                            Alfatec-Pharma GmbH, Germany
  SOURCE:
                            Ger. Offen., 6 pp.
                            CODEN: GWXXBX
  DOCUMENT TYPE:
                            Patent
  LANGUAGE:
                            German
  FAMILY ACC. NUM. COUNT:
  PATENT INFORMATION:
       PATENT NO.
                        KIND DATE
                                               APPLICATION NO. DATE
       DE 4140177
                         A1
                               19930609
                                               DE 1991-4140177 19911205
       DE 4140177
                          C2
                               19951221
       WO 9310767
                         A1
                               19930610
                                               WO 1992-DE1009
                                                                 19921204
          W: AU, CA, JP, US
          RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
      WO 9310768
                         A1
                              19930610
                                               WO 1992-DE1010 19921204
          W: AU, CA, JP, US
          RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
      AU 9230801
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                                               AU 1992-30801
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      AU 671964
                         B2
                              19960919
      AU 9230802
                         A1
B2
                              19930628
                                               AU 1992-30802
                                                                 19921204
      AU 671965
                              19960919
      EP 615444
                         A1
                             19940921
                                               EP 1992-924546
                                                                19921204
      EP 615444
                         B1
                              19960306
          R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, NL, PT, SE
      EP 615445
                        A1 19940921
                                              EP 1992-924547 19921204
      EP 615445
                         B1
                              19960515
          R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, NL, PT, SE
      AT 134875
                         Ε
                              19960315
                                              AT 1992-924546 19921204
      ES 2085656
                         Т3
                              19960601
                                               ES 1992-924546
                                                                19921204
      AT 137962
                         Ε
                              19960615
                                              AT 1992-924547
                                                                19921204
      ES 2087565
                         T3
                              19960716
                                              ES 1992-924547
                                                                19921204
      US 5614219
                         Α
                              19970325
                                              US 1994-244691
                                                                 19940913
      US 5932245
                         Α
                              19990803
                                              US 1994-244615
                                                                19941025
 PRIORITY APPLN. INFO.:
                                           DE 1991-4140177 A 19911205
DE 1991-4140178 A 19911205
                                           DE 1991-4140186 A 19911205
DE 1991-4140195 A 19911205
                                           US 1992-876867 A 19920430
                                           WO 1992-DE1009 A 19921204
WO 1992-DE1010 A 19921204
     A rapid-release antidiabetic nanosol formulation comprises a
     glibenclamide (I) inner phase (10-800 nm particle size) and a gelatin or
     collagen hydrolyzate outer phase. The two phases have opposite isoionic elec. charges. A soln. (pH 2.2; HCl) of 500 g gelatin B in 3L water was blended with a soln. of 13.89 g I in 0.2 L EtOH to give a nanosol
       EtOH was evapd. in vacuum, followed by dry-spraying and tabletting of
     the product.
L10 ANSWER 47 OF 47 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER:
                          1992:654275 CAPLUS
DOCUMENT NUMBER:
                          117:254275
TITLE:
                          Nanostructures in sol-gel derived materials:
                          application to the elaboration of nanofiltration
                          membranes
AUTHOR (S):
                          Guizard, C.; Julbe, A.; Larbot, A.; Cot, L.
CORPORATE SOURCE:
                          Lab. Physicochim. Mater., Ec. Norm., Montpellier,
                          34053, Fr.
SOURCE:
                          Journal of Alloys and Compounds (1992), 188, 8-13
                          CODEN: JALCEU; ISSN: 0925-8388
DOCUMENT TYPE:
                          Journal; General Review
                          English
    A review with 17 refs. Development of membranes resistant to heating and
     chems. is expected for the sepn. of small mols. in biotechnologies,
    pharmaceutics, chem. industries, water treatment, and also in gas sepn.
    When aiming at sepn. of small mols. (mol. wt. <1000) or multivalent ions,
    nanoscale pores or almost dense materials will be required. Pure inorg.
    materials as well as org./inorg. polymers are very good candidates for
    prepg. this new generation of membranes. The nanostructural organization
    of sol-gel-derived materials means that the sieve concept developed for
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microfiltration (MF) and ultrafiltration (UF) membranes can be surpassed and a concept of intelligent membrane can be recognized for nanofiltration

membranes. In fact, specific interactions are expected between the nanophased membranes and solubilized species or gas mols. passing through the membrane. Provided that these interactions can be managed, an increase in membrane performances is obtained, with the possibility of developing new application fields. Examples are given showing the important role played by the basic properties of ceramics and organically modified ceramics in the prodn. of tailored nanofiltration membranes.